

STIC-ILL

vol w 437391

From: STIC-Biotech/ChemLib
Sent: Thursday, March 20, 2003 7:20 AM
To: STIC-ILL
Subject: FW: 09850697

-----Original Message-----

From: Yaen, Christopher
Sent: Wednesday, March 19, 2003 5:00 PM
T : STIC-Biotech/ChemLib
Subject: 09850697

10027135

could you please get the following ref(s):

Thromb Haemost 1989 Nov 24;62(3):846-9

2595658

Appl Environ Microbiol 1994 Aug;60(8):2793-801

Antonie Van Leeuwenhoek 1996 Feb;69(2):151-9

Oncogene 2000 Mar 16;19(12):1579-88

Christopher Yaen
Patent Examiner
US PTO
Art Unit 1642
CM1-Rm 8E18
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703-305-3586

STIC-ILL

NO

437370

From: STIC-Biotech/ChemLib
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To: STIC-ILL
Subject: FW: 09850697

-----Original Message-----

From: Yaen, Christopher
Sent: Wednesday, March 19, 2003 5:00 PM
To: STIC-Biotech/ChemLib
Subject: 09850697

10026554

could you please get the following ref(s):

Thromb Haemost 1989 Nov 24;62(3):846-9

Appl Environ Microbiol 1994 Aug;60(8):2793-801

Antonie Van Leeuwenhoek 1996 Feb;69(2):151-9

8775975

Oncogene 2000 Mar 16;19(12):1579-88

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NEWS 38 Dec 30 ISMEC no longer available
NEWS 39 Jan 13 Indexing added to some pre-1967 records in CA/CAPLUS
NEWS 40 Jan 21 NUTRACEUT offering one free connect hour in February 2003
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NEWS 42 Simultaneous left and right truncation added to COMPENDEX,
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NEWS 43 Feb 13
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 YAO, Monique, G., 1189 Woodgate Drive, Carmel, IN 46033, US [US, US];
 YUE, Henry, 826 Lois Avenue, Sunnyvale, CA 94087, US [US, US];
 YUE, Huibin, 1170 South Stelling Road, Cupertino, CA 95014, US [US, US];

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 IN

PROCESSING COMPLETED FOR L2
 L3 12 DUP REM L2 (1 DUPLICATE REMOVED)

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CA 94304, US
English
LA
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Patent
PI
WO 2003014322 A2 20030220
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NO NZ OM PH PL PT RO RU SD SE SG SI SK SL TJ TM TN TR TT TZ
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RW (EAP): AM AZ BY KG KZ MD RU TJ TM
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AI WO 2002-0525465 A 20020808
PRAI US 2001-60/311.017 20010808
US 2001-60/313.070 20010817
US 2001-60/313.071 20010817
US 2001-60/314.678 20010824
US 2001-60/316.692 20010831
US 2001-60/317.913 20010907
US 2001-60/322.182 20010914
US 2001-60/340.747 20011207
US 2001-60/342.761 20011220
US 2002-60/369.129 20020329
L3 ANSWER 2 OF 12 CAPLUS COPYRIGHT 2003 ACS DUPLICATE 1
DN 2002:755245 CAPLUS
DN 137:274175
TI DNA, CDNA and protein sequences of spermatogenesis assocd. factors a
method for diagnosis of cancer
IN Kulesz-Martin, Molly F.; Liu, Yungang
PA USA
U.S. Pat. Appl. Publ., 42 pp., Cont.-in-part of U.S. Ser. No. 777,753.
CODEN: USXXCO
DT Patent
LA English
FAN CNT 1
PATENT NO. KIND DATE APPLICATION NO. DATE

PI US 2002143169 AI 20021003
PRAI US 1997-938308 B2 19970927 US 2001-850697 20010508
US 2001-777753 A2 20010206
L3 ANSWER 3 OF 12 USPATFULL
AN 2002:301167 USPATFULL
TI Nucleic acids, proteins, and antibodies
IN Rosen, Craig A., Laytonville, MD, UNITED STATES
Ruben, Steven N., Olney, MD, UNITED STATES
Barash, Steven C., Rockville, MD, UNITED STATES
PI US 2002168711 AI 20021114
AI US 2001-764868 AI 20010117 (9)
PRAI US 2000-179065P 20000131 (60)
US 2000-180628P 20000204 (60)
US 2000-214886P 20000628 (60)
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US 2000-237037P 20001002 (60)
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US 2000-240960P 20001020 (60)
US 2000-239935P 20001013 (60)
DT Utility
FS APPLICATION
LN CNT 31967
INCL INCLM: 435/069.100
INCLM: 435/320.100; 435/183.000; 530/350.000; 536/023.100
NCL INCLM: 435/069.100
NCLM: 435/325.000; 435/320.100; 435/183.000; 530/350.000; 536/023.100
IC [7]
ICM: C12P021-02
ICS: C12N005-06; C07H021-04; C12N009-00; C07K014-435
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L3 ANSWER 4 OF 12 USPATFULL
AN 2002:112541 USPATFULL
TI Proteins related to schizophrenia and uses thereof
IN St. George-Hyslop, Peter H., Toronto, CANADA
Fraser, Paul E., Toronto, CANADA
PA The Governing Council of the University of Toronto (non-U.S. corporation)
PI US 2002058276 AI 20020516
AI US 2001-945258 AI 20010831 (9)
PRAI US 2000-229889P 20000901 (60)
DT Utility
FS APPLICATION
LN CNT 2909
INCL INCLM: 435/006.000
INCLM: 424/009.200; 800/003.000
NCL INCLM: 435/006.000

IC NCLS: 424/009.200: 800/003.000

AN ICM: C12Q001-68

ICS: A61K049-00: A01K067-00

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L3 ANSWER 5 OF 12 PCTFULL COPYRIGHT 2003 Univentio
AN 2002099062 PCTFULL ED 20021218 EW 200250

T1EN NOVEL ANTIBODIES THAT BIND TO ANTIGENIC POLYPEPTIDES, NUCLEIC ACIDS

ENCODING THE ANTIGENS, AND METHODS OF USE

T1FR NOUVEAUX ANTICORPS SE FIXANT A DES POLYPEPTIDES ANTIGENIQUES, ACIDES

NUCLEIQUES CODANT LES ANTIGENES ET MODES D'UTILISATION

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US], for US only;

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US only

ELRIFI, Ivor, R., Mintz, Levin, Cohn, Ferris, Glovsky, and Popeo, P.,

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[CN, US]

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RW (OAPI): BF BJ CF CG CI CM CA GN GQ GW ML MR NE SN TD TG
WO 2002-017559 A 20020604
US 2001-60/295,607
US 2001-60/296,418 20010606
US 2001-60/296,404 20010606
US 2001-60/296,575 20010607
US 2001-60/297,414 20010611
US 2001-60/297,573 20010612
US 2001-60/297,567 20010612
US 2001-60/298,285 20010614
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US 2002-60/371,523 20020410
US 2002-60/371,346 20020410
US 2002-10/161,493 20020603
L3 ANSWER 6 OF 12 PCTFULL COPYRIGHT 2003 Univentio
AN 2002046395 PCTFULL ED 20020624 EW 200224
TIEN ENZYMES
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TANG, Y., Tom, 4230 Ranwick Court, San Jose, CA 95118, US [US, US];
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BAUGHN, Mariah, R., 14244 Santiago Road, San Leandro, CA 94577, US [US, US];
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LEE, Sally, 825 East Evelyn, #425, Sunnyvale, CA 94086, US [US, US], for US only;
RAMKUMAR, Jayalaxmi, 34359 Maybird Circle, Fremont, CA 94555, US [IN, US], for US only;
WARREN, Bridget, A., 10130 Parkwood Drive #2, Cupertino, CA 95014, US [US, US], for US only;
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LAF English
LA Patent
DT WO 2002046395
PI AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ
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UZ VN YU ZA ZW
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AI WO 2001-US47432 A 20011204
PRAI US 2000-60/251,824 20001207
US 2000-60/254,312 20001208
US 2000-60/255,773 20001214
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US 2000-60/257,488 20001221
US 2001-60/262,839 20010119
US 2001-60/264,402 20010126
ICM C12N009-00

US], for all designates States except US;
TANG, Y., Tom, 4230 Ranwick Court, San Jose, CA 95118, US [US, US], for US only;
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HILLMAN, Jennifer, L., 230 Monroe Drive, #17, Mountain View, CA 94040, US [US, US], for US only;
HAMLET-COX, Diana, Incyte Genomics, Inc., 3160 Porter Drive, Palo Alto, CA 94304, US [US, US], for US only;
LAF English
LA Patent
DT WO 2002046395
PI AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ
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RW (OAPI): BF BJ CF CG CI CM GA GN GQ GW ML MR NE SN TD TG
AI WO 2001-US47432 A 20011204
PRAI US 2000-60/251,824 20001207
US 2000-60/254,312 20001208
US 2000-60/255,773 20001214
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US 2001-60/264,402 20010126
ICM C12N009-00

L3 ANSWER 7 OF 12 PCTFULL COPYRIGHT 2003 Univentio
AN 2002029113 PCTFULL ED 20020627 EW 200215
TIEN METHODS FOR MONITORING MULTIPLE GENE EXPRESSION
TIFR METHODES DE SURVEILLANCE DE L'EXPRESSION GENETIQUE MULTIPLE
IN BERKA, Randy, 3609 Modoc, Davis, CA 95616, US;
CLAUSEN, Ib, Groth, Fyrestein 6, DK-3400 Hillerod, DK
PA NOVOZYMES BIOTECH, INC., 1445 Drew Avenue, Davis, CA 95616, US (US, US);
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AG STARNES, Robert, Novozymes Biotech, Inc., 1445 Drew Avenue, Davis, CA
95616, US
LAF English
LA Patent
DT WO 2002029113 A2 20020411
PI AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ
DS DE DK DM DZ EC EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP
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UZ VN YU ZA ZW
RW (ARIPO): GH GM KE LS MW MZ SD SL SZ TZ UG ZW
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AI WO 2001-US31437 A 20011005
PRAI US 2000-09/680,598 20001006
US 2001-60/279,526 20010327
ICM C12Q001-68

L3 ANSWER 8 OF 12 PCTFULL COPYRIGHT 2003 Univentio
AN 2002018434 PCTFULL ED 20020705 EW 200210
TIEN PROTEINS RELATED TO SCHIZOPHRENIA AND USES THEREOF
TIFR PROTEINES LIEES A LA SCHIZOPHRENIE ET UTILISATIONS DE CELLULES-CI
IN ST.GEORGE-HYSLAP, Peter, H., 210 Richview Avenue, Toronto, Ontario M5P
3G3, CA;
FRASER, Paul, E., 611 Windermere Avenue, Toronto, Ontario M6S 3L9, CA
PA THE GOVERNING COUNCIL OF THE UNIVERSITY OF TORONTO, Sincove Hall, Room
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LAF English
LA Patent
DT WO 2002018434 A2 20020307
PI AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ
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NO NZ PH PL PT RO RU SD SE SG SI SK SL TJ TM TR TT TZ UA UG
UZ VN YU ZA ZW
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AI WO 2001-CA1243 A 20010831
PRAI US 2000-60/229,889 20000901
ICM C07K014-47

L3 ANSWER 9 OF 12 PCTFULL COPYRIGHT 2003 Univentio
AN 200006339 PCTFULL ED 20020814
TIEN NOVEL PROTEINS AND NUCLEIC ACIDS ENCODING SAME
TIFR NOUVELLES PROTEINES ET ACIDES NUCLEIQUES LES CODANT
IN SPADERNA, Steven, K.;
TCHERNEV, Velizar;
LIU, Xiaohong;
SHENOY, Suresh;
SPYTEK, Kimberly;

L3 ANSWER 10 OF 12 PCTFULL COPYRIGHT 2003 Univentio
AN 2001054733 PCTFULL ED 20020827
TIEN NUCLEIC ACIDS, PROTEINS AND ANTIBODIES
TIFR ACIDES NUCLEIQUES, PROTEINES ET ANTICORPS
IN ROSEN, Craig, A.;
BARASH, Steven, C.;
RUBEN, Steven, M.
HUMAN GENOME SCIENCES, INC.;
ROSEN, Craig, A.;
BARASH, Steven, C.;
RUBEN, Steven, M.

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PATTURAJAN, Meera;
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SZEKERES, Edward, S.;
ALSOBROOK, John, II;
LEPLEY, Denise, M.;
SHEN, Lei;
BURGESS, Catherine, E.;
SHIMKETS, Richard, A.;
PADIGARU, Muralidhara
CURAGEN CORPORATION;
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TCHERNEV, Velizar;
LIU, Xiaohong;
SHENOY, Suresh;
SPYTEK, Kimberly;
ZERHUSEN, Bryan;
PATTURAJAN, Meera;
TAUPIER, Raymond, J.;
RASTELLI, Luca;
GROSSE, William, M.;
SZEKERES, Edward, S.;
ALSOBROOK, John, II;
LEPLEY, Denise, M.;
SHEN, Lei;
BURGESS, Catherine, E.;
SHIMKETS, Richard, A.;
PADIGARU, Muralidhara
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DT WO 2002006339 A2 20020124
PI AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ
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NL PT SE TR BF BJ CF CG CI CM GA GN GW ML MR NE SN TD TG
AI WO 2001-US21249 A 20010703
PRAI US 2000-60/215,854 20000703
US 2000-60/215,856 20000703
US 2000-60/215,902 20000703
US 2000-60/216,585 20000707
US 2000-60/216,586 20000707
US 2000-60/216,722 20000707
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US 2000-60/218,992 20000717
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US 2001-60/268,734 20010214
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ICM C07K014-47

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US 2000-60/249, 209 20001117
US 2000-60/249, 300 20001117
US 2000-60/249, 265 20001117
US 2000-60/250, 391 20001201
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US 2000-60/256, 719 20001205
US 2000-60/251, 030 20001205
US 2000-60/251, 988 20001205
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US 2000-60/251, 869 20001208
US 2000-60/251, 856 20001208
US 2000-60/251, 868 20001208
US 2000-60/251, 990 20001208
US 2000-60/251, 989 20001208
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US 2000-60/259, 678 200010105
A61K048-00
C12N005-00

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ANSWER 11 OF 12 PCTFULL COPYRIGHT 2003 Univentio
2001045007 PCTFULL ED 20020827
A METHOD AND SYSTEM FOR DISCOVERY OF TRADES BETWEEN PARTIES
PROCEDE ET SYSTEME PERMETTANT DE DECOUVRIR DES MARCHES ENTRE DES PARTIES
MACREADY, William, G.;
EL-BELTAGY, Mohammed;
ROY, Barbeau;
ANDERSON, Mark
BIOS GROUP INC.
Patent
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AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CR CU CZ DE
DK DM DZ EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG
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WO 2000-US33017 A 20001206
US 1999-60/168, 754 19991206
US 2000-60/194, 880 20000406
G06F017-60

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ANSWER 12 OF 12 PCTFULL COPYRIGHT 2003 Univentio
2000060069 PCTFULL ED 20020515
A PRESENILIN ASSOCIATED MEMBRANE PROTEIN AND USES THEREOF
PROTEINE MEMBRANAIRE ASSOCIEE A LA PRESENILINE ET SES UTILISATIONS
ST. GEORGE-HYSLOP, Peter, H.;
FRASER, Paul, E.
THE GOVERNING COUNCIL OF THE UNIVERSITY OF TORONTO
English
Patent
W:

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LA
DT
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AE AG AL AM AT AU AZ BA BB BG BR BY CA CH CN CR CU CZ DE DK
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AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE BF BJ
CF CG CI CM CA CN CH GM ML MR NE SN TD TG

WO 2000-CA354
US 1999-60/127, 452 19990401
US 1999-60/173, 826 19991230
C12N015-12
C07K014-705; A01K067-027; C12N005-10; C12Q001-68; G01N033-50

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L3 ANSWER 12 OF 12 PCTFULL COPYRIGHT 2003 Univentio

DETD Furthermore, mutant PAMP nucleic acids, proteins, or peptides, cells transfected with vectors comprising mutant PAMP nucleic acids, transgenic animals expressing mutant PAMP or peptides thereof, and their use in studying Alzheimer's Disease and other.

The invention also provides vectors, and particularly expression vectors (e.g., cos-Tet vector), which include any of the above-described SUBSTITUTE SHEET (RULE 26) nucleic acids. It is a further object of the invention to provide vectors in which normal or mutant PAMP nucleic acid sequences are operably joined to exogenous regulatory regions to produce altered patterns of expression.

phenotype on the cell in which it is expressed. The term expression system means a host cell transformed by a compatible expression vector and cultured under suitable conditions e.g. for the expression of a protein coded for by foreign DNA carried by the vector and introduced to the host cell.

The terms vector, cloning vector and expression vector mean the vehicle by which a DNA or RNA sequence (e.g., a foreign gene) can be introduced into a host cell, so.

Vectors include plasmids, phages, viruses, etc. A cassette refers to a DNA coding sequence or segment of DNA that codes for an expression product that can be inserted into a vector at defined restriction sites. The cassette restriction sites are designed to ensure insertion of the cassette in the proper reading frame. Generally, foreign DNA is inserted at one or more sites of the vector DNA, and then is carried by the vector into a host cell along with the transmissible vector DNA. A segment or sequence of DNA having inserted or added DNA, such as an expression vector, can also be called a DNA construct. Recombinant cloning vectors will often include one or more SUBSTITUTE SHEET (RULE 26) replication systems for cloning or expression, one or more markers for selection in.

In the context of the present invention, an gene is heterologous to the recombinant vector DNA in which it is inserted for cloning or expression, and it is heterologous to a host cell containing such a vector, in which it is expressed, e.g., a CHO cell.

via Omay

pumps inserted into the cerebral ventricles); via the transplantation of genetically-modified cells expressing recombinant genes; or via the use of biological vectors (e.g., retrovirus, adenovirus, adeno-associated virus, lentivirus, or herpes simplex virus-based vectors) which allow targeted expression of appropriately modified gene products in selected cell types. It should be noted that the recombinant proteins described. . . or part of PAMP or PAMP mutants, e.g., as mini-gene cDNA transgene constructs under the regulation of suitable promoter elements carried in vectors such as cos-Tet for transgenic mice and pcDNA (Invitrogen, California) in transfected cell lines. . . to identify upstream and downstream modifiers of a PAMP phenotype. Transgenic animals can also be prepared by introducing the transgene on a vector; such animals, which are not modified in the germ line and are only transiently transgenic, naturally cannot pass along the genetic information. . . may be truncated), 1.5 can be introduced in vivo, ex vivo, or in vitro using a viral or a non-viral vector, e.g., as discussed above. Expression in targeted tissues can be effected by targeting the transgenic vector to specific cells, such as with a viral vector or a receptor ligand, or by using a tissue-specific promoter, or both. Targeted gene delivery is described in International Patent Publication WO. . . Preferably, for in vivo administration, an appropriate immunosuppressive treatment is employed in conjunction with the viral vector, e.g., adenovirus vector, to avoid immuno-deactivation of the viral vector and transduced cells. For example, immunosuppressive cytokines, such as interleukin 12 (IL-12), interferon- γ (IFN γ), or anti-CD4 antibody, can be administered to block humoral or cellular immune responses to the viral vectors (see, e.g., Wilson, Nature Medicine, 1995). In that regard, it is advantageous to employ a viral vector that is engineered to express a minimal number of antigens.

Herpes virus vectors. Because herpes virus is trophic for cells of the nervous system (neural cells), it is an attractive vector for delivery of function PAMP genes. Various defective (non-replicating, and thus non-infectious) herpes virus vectors have been described, such as a defective herpes virus 1 (HSV1) vector (Kaplit et al., Molec. Cell. Neurosci. 2:320-330, 1991, International Patent Publication No. WO 94/21807, published SUBSTITUTION SHEET (RULE 26) September 29, 1994; International. . .

Adenovirus vectors. Adenoviruses are eukaryotic DNA viruses that can be modified to efficiently deliver a nucleic acid of the invention to a . . . adenovirus, more preferably a CAV2 adenovirus (e.g., Mahattan or A26/61 strain (ATCC VR-800), for example). Various replication defective adenovirus and minimum adenovirus vectors have been described for gene therapy (WO94126914, WO95/026971 WO94/28938, SUBSTITUTION SHEET (RULE 26) which is infected with a human helper virus (for. . . Retrovirus vectors. In another embodiment the gene can be introduced in a retroviral vector, e.g., as described in Anderson et al., U.S.

infect dividing cells. The retrovirus genome includes two LTRs, an encapsidation sequence and three coding regions (gag, pol and env). In recombinant retroviral vectors, the gag, pol and env genes are generally deleted, in whole or in part, and replaced with a heterologous nucleic acid. . . These vectors can be constructed from different types of retrovirus, such as MoMuLV (murine Moloney leukemia virus), MEV (murine Moloney sarcoma virus), HaSV (Harvey. . . line PA317 (US 4,861,719); the PsiCRIP cell line (WO 90/02806) and the GP+envAm-12 cell line (WO 89/07150). In addition, the recombinant retroviral vectors can contain modifications within the LTRs for suppressing transcriptional activity as well as extensive encapsidation sequences which may include a part of the gag gene (Bender et al., J. Virol. 61:1639, 1987). Recombinant retroviral vectors are purified by standard techniques known to those having ordinary skill in the art.

Retrovirus vectors can also be introduced by recombinant DNA viruses, which permits one cycle of retroviral replication and amplifies transfection efficiency (see WO 95/22617, . . .

Lentivirus vectors. In another embodiment, lentiviral vectors are can be used as agents for the direct delivery and sustained expression of a transgene in several tissue types, including brain, retina, muscle, liver and blood. The vectors can efficiently transduce dividing and non-dividing cells in SUBSTITUTION SHEET (RULE 26) these tissues, and maintain long-term expression of the gene of. . . 72,9873-801,1998). Lentiviral packaging cell lines are available and known generally in the art. They facilitate the production of high-titer lentivirus vectors for gene therapy. An example is a tetracycline-inducible VSV-G pseudotyped lentivirus packaging cell line which can generate virus particles at titers greater than 10⁶ IU/ml for at least 3 to 4 days (Kafri, et al., J. Virol., 73: 576-584, 1999). The vector

produced by the inducible cell line can be concentrated as needed for efficiently transducing nondividing cells in vitro and in vivo.

Non-viral vectors. A vector can be introduced in vivo in a non-viral vector, e.g., by lipofection, with other transfection facilitating agents (peptides, polymers, etc.), or as naked DNA. Synthetic cationic lipids can be used to . . .

DNA vectors for gene therapy can be introduced into the desired host cells by methods known in the art, e.g., electroporation, microinjection, cell fusion, DEAE dextran, calcium phosphate precipitation, use of a gene gun (ballistic transfection), or use of a DNA vector transporter (see, e.g., Wu et al., J. Biol. . .

either V5-tagged PAMP or empty plasmid (mock transfection control). Duplicate experiments were performed by: (1) transient transfection of V5-PAMP and PAPP695 (or empty vector plus PAPP695 as a mock transfection control) into murine embryonic fibroblasts stably infected with human PSI expressed from a retroviral vector construct (Clontech, CA) - or (2) transient transfection of V5-PAMP (or empty plasmid) into HEK293 cell lines with a stable expression of. . .

Cells were transiently transfected with PAMP cDNA (SEQ ID NO: 13) tagged at the 3'-end with a V5-epitope encoded from the pCDNA6 vector. The conditioned media were collected 20 hr after transient transfection with PAMP (or with empty vector), and the Ap40 and Ap42 levels were measured by ELISA (Zhang L, et al., J Biol Chem 1999; 274: 8966). In. . .

region D169L. PAMR0312-369 in the central conserved region D140X: PAMPA312-340 in the central conserved region YDT:- PAMPD458A in the putative 'aspartyl protease' DTA site SPAP- PAMPF633A/F635A in the SPAP motif TM: PAMPS683A in the TM domain C31D: PAMPA630-668 in the conserved region adjacent to the TM domain To further examine the role of. . . above mutations, as well as normal/wild type PAMP (PAMPwt) ONA and the ONA for an unrelated protein (LacZ), in frame into pCDNA6 vectors. A series of HEK293 cell lines stably expressing endogenous PSI 7 PAPPswedish and either wild type PAMP or PAMP constructs in which. . .

or in the AP42/AK ratio, when the PAMPwt, PAMPD458A, PAMPA630-668, PAMPF633A/F635A, and PAMPS683A cells were compared to control lines (expressing LacZ, or

empty vector).

Mock (LacZ/empty 1.0 1.0 1.0 vector)

Wild type nicastrin 1.03 \pm plum; 0.09 1.05 \pm plum; 0.07 0.99 \pm plum; 0.07 0.07 D336A/V337A 3.09 \pm plum; 0.50 1.61 \pm plum; 0.19 1.81 \pm plum; 0.15 (p. . .

CLMEN 10 A vector comprising the nucleic acid of claim 7 operatively associated with an expression control sequence.

11 A cell transfected with the vector of claim 1 0.

15 A vector comprising the nucleic acid of claim 13 operatively associated with an expression control sequence.

0 16. A cell transfected with the vector of claim 15.

=> d kwic 1

L3 ANSWER 1 OF 12 FCTFULL COPYRIGHT 2003 Univentio ABEN . . . cell growth, differentiation, and death (CGDD) and polynucleotides which identify and encode CGDD. Embodiments of the invention also provide expression vectors, host cells, antibodies, agonists, and antagonists. Other embodiments provide methods for diagnosing, treating, or preventing disorders associated with aberrant expression. . .

DETD . . . Hamada, K. et al. (1996; Cancer Res. 56:3047-3054) are investigating the introduction of p53 into cervical cancer cells via an adenoviral vector as an experimental therapy for cervical cancer.

Spermatogenesis associated factor (SPAF) is an AAA-protein (Afpase associated with diverse activities) specific to early spermatogenesis and malignant conversion. SPAF is expressed in spermatogonia and early spermatocytes in the basal compartment of the seminiferous tubules (Liu, Y.

Table 6 provides an appendix which describes the tissues and vectors used for construction of the cDNA libraries shown in Table 5.

are now described. All publications mentioned herein are cited for the purpose of describing and disclosing the cell lines, protocols, reagents and vectors which are reported in the publications and which might be used in connection with various embodiments of the invention. Nothing herein. . .

include a nucleic acid sequence operably linked to a promoter sequence. Such a recombinant nucleic acid may be part of a vector that is used, for example, to transform a cell.

Alternatively, such recombinant nucleic acids may be part of a viral vector, e.g., based on a vaccinia virus, that could be used to vaccinate a mammal wherein the recombinant nucleic acid is expressed, inducing. . .

or by infection with a recombinant virus. In another embodiment, the nucleic acid can be introduced by infection with a recombinant viral vector, such as a lentiviral vector (Lois, C. et al. (2002) Science 295:868-872). The term genetic manipulation does not include classical cross-breeding, or in vitro fertilization, but rather.

is 53% identical from residue M1 to residue K56, and 84% identical from residue G55 to residue G535, to Mus musculus SPAP (spermatogenesis associated factor, AAA family) (GenBank ID g4105619) as determined by the Basic Local Alignment Search Tool (BLAST). (See Table 2.) The.

most frequently represented by the Incyte cDNA sequences which were used to assemble and confirm the above polynucleotides. The tissues and vectors which were used to construct the cDNA libraries shown in Table 5 are described in Table 6. fragments thereof, entirely by synthetic chemistry. After production, the synthetic polynucleotide may be inserted into any of the many available expression vectors and cell systems using reagents well known in the art. Moreover, synthetic chemistry may be used to introduce mutations into a polynucleotide.

method which may be employed, restriction-site PCR, uses universal and nested primers to amplify unknown sequence from genomic DNA within a cloning vector (Sarkar, G. (1993) PCR Methods Appl. 2:318-322). Another method, inverse PCR, uses primers that extend in divergent directions to amplify unknown.

order to express a biologically active CGDD, the polynucleotides encoding CGDD or derivatives thereof may be inserted into an appropriate expression vector, i.e., a vector which contains the necessary elements for transcriptional and translational control of the inserted coding sequence in a suitable host. These elements include regulatory sequences, such as enhancers, constitutive and inducible promoters, and 5' and 3' untranslated regions in the vector and in polynucleotides encoding CGDD. Such elements may vary in their strength and specificity. Specific initiation signals may also be used to. . . in cases where a polynucleotide sequence encoding CGDD and its initiation codon and upstream regulatory sequences are inserted into the appropriate expression vector, no additional transcriptional or translational control signals may be needed. However, in cases where only coding sequence, or a fragment thereof, is inserted, exogenous translational control signals including an in-frame ATG initiation codon should be provided by the vector. Exogenous translational elements and initiation codons may be of various origins, both natural and synthetic. The efficiency of expression may be.

Methods which are well known to those skilled in the art may be used to

construct expression

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vectors containing polynucleotides encoding CGDD and appropriate transcriptional and translational control elements. These methods include in vitro recombinant DNA techniques, synthetic techniques, and.

A variety of expression vector/host systems may be utilized to contain and express polynucleotides encoding CGDD. These include, but are not limited to, microorganisms such as bacteria transformed with recombinant bacteriophage, plasmid, or cosmid DNA expression vectors; yeast transformed with yeast expression vectors; systems infected with viral expression vectors (e.g., baculovirus); plant cell systems transformed with viral expression vectors (e.g., cauliflower mosaic virus, CaMV, or tobacco mosaic virus, TMV) or with bacterial expression vectors (e.g., Ti or pBR322 plasmids); or animal cell systems (Sambrook, supra; Ausubel et al., supra; Van Heeke, G. and S.M. Schuster (1989)).

Sci. USA 81:3655-3659; Harrington, J.J. et al. (1997) Nat. Genet. 15:345-355). Expression vectors derived from retroviruses, adenoviruses, or herpes or vaccinia viruses, or from various bacterial plasmids, may be used for delivery of polynucleotides to.

In bacterial systems, a number of cloning and expression vectors may be selected depending upon the use intended for polynucleotides encoding CGDD. For example, routine cloning, subcloning, and propagation of polynucleotides encoding CGDD can be achieved using a multifunctional E. coli vector such as PBLUESCRIPT (Stratagene, La Jolla CA) or PSORT1 plasmid (Invitrogen). Ligation of polynucleotides encoding CGDD into the vector's multiple cloning site disrupts the lacZ gene, allowing a colorimetric screening procedure for identification of transformed bacteria containing recombinant molecules. In addition, these vectors may be useful for in vitro transcription, dideoxy sequencing, single strand rescue with helper phage, and creation of nested deletions in the.

264:5503-5509). When large quantities of CGDD are needed, e.g. for the production of antibodies, vectors which direct high level expression of CGDD may be used. For example, vectors containing

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the strong, inducible SP6 or T7 bacteriophage promoter may be used. Yeast expression systems may be used for production of CGDD. A number of vectors containing constitutive or inducible promoters, such as alpha factor, alcohol oxidase, and PGH promoters, may be used in the yeast Saccharomyces cerevisiae or Pichia pastoris. In addition, such vectors direct either the secretion or intracellular retention of expressed proteins and enable integration of foreign polynucleotide sequences into the host genome.

complement of the polynucleotide encoding CGDD may be administered to a subject to treat or prevent a disorder associated. . . .

In other embodiments, any protein, agonist, antagonist, antibody, complementary sequence, or vector embodiment may be administered in combination with other appropriate therapeutic agents. . . .

Antisense sequences can also be introduced intracellularly through the use of viral vectors, such as retrovirus and adeno-associated virus vectors (Miller, A.D. (1990) Blood 76:27-1; Ausubel et al., supra; Uckert, W. and W. Walther (1994) Phannacol. Ther. 63:323-347). Other. . . .

In a further embodiment of the invention, diseases or disorders caused by deficiencies in CGDD are treated by constructing mammalian expression vectors encoding CGDD and introducing these vectors by mechanical means into CGDD-deficient cells. Mechanical transfer technologies for use with cells in vivo or ex vitro include (i) direct. . . .

Expression vectors that may be effective for the expression of CGDD include, but are not limited to, the PCDNA 3.1, EPITAG, PRCMV2, PREP, PVAX, PCR2-TOPOTA vectors (Invitrogen, Carlsbad CA), PCMV-SCRIPT, PCMV-TAG, PEGSH/PERV (Stratagene, La Jolla CA), and PTET-OFF, PTET-CN, PTRE2, PTRE2-LUC, PTK-HYG (Clontech, Palo Alto CA). CGDD may be. . . .

of the invention, diseases or disorders caused by genetic defects with respect to CGDD expression are treated by constructing a retrovirus vector consisting of (i) the . . .

polynucleotide encoding CGDD under the control of an independent promoter or the retrovirus long terminal repeat (LTR). . . . RNA packaging signals, and (iii) a Rev-responsive element (RRE) along with additional retrovirus cis-acting RNA sequences and coding sequences required for efficient vector propagation. Retrovirus vectors (e.g., PB and PRENeo) are commercially available (Stratagene) and are based on published data (Riviere, I. et al. (1995) Proc. Natl. Acad. Sci. USA 92:6733-6737), incorporated by reference herein. The vector is propagated in an appropriate vector producing cell line (VPCL) that expresses an envelope gene with a tropism for receptors on the target cells or a promiscuous. . . .

transducing efficiency retroviral supernatant) discloses a method for obtaining retrovirus packaging cell lines and is hereby incorporated by reference. Propagation of retrovirus vectors, transduction of a population of cells (e.g., CD4⁺ T-cells), and the return of transduced cells to a patient are procedures. . . .

cells which have one or more genetic abnormalities with respect to the expression of CGDD. The construction and packaging of

mammalian cells, a number of viral-based expression systems may be utilized. In cases where an adenovirus is used as an expression vector, polynucleotides encoding CGDD may be ligated into an adenovirus transcription/translation complex consisting of the late promoter and tripartite leader sequence. Insertion in. . . .

enhancers, such as the Rous sarcoma virus (RSV) enhancer, may be used to increase expression in mammalian host cells. SV40 or EBV-based vectors may also be used for high-level protein expression. . . .

stable expression of CGDD in cell lines is preferred. For example, polynucleotides encoding CGDD can be transformed into cell lines using expression vectors which may contain viral origins of replication and/or endogenous expression elements and a selectable marker gene on the same or on a separate vector. . . .

Following the introduction of the vector, cells may be allowed to grow for about 1 to 2 days in enriched media before being switched to selective media. . . .

not only to identify transformants, but also to quantify the amount of transient or stable protein expression attributable to a specific vector system (Rhodes, C.A. (1995) Methods Mol. Biol. 55:121-131). . . .

Alternatively, polynucleotides encoding CGDD, or any fragments thereof, may be cloned into a vector for the production of an mRNA probe. Such vectors are known in the art, are commercially available, and may be used to synthesize RNA probes in vitro by addition of. . . .

from cell culture. The protein produced by a transformed cell may be secreted or retained intracellularly depending on the sequence and/or the vector used. As will be understood by those of skill in the art, expression vectors containing polynucleotides which encode CGDD may be designed to contain signal sequences which direct secretion of CGDD through a prokaryotic or eukaryotic. . . .

129/SvJ cell line, are derived from the early mouse embryo and grown in culture. The ES cells are transformed with a vector containing the gene of interest disrupted by a marker gene, e.g., the neomycin phosphotransferase gene (neo; Capecchi, M.R. (1989) Science 244:1288-1292). The vector integrates into the corresponding region of the host genome by homologous recombination. Alternatively, homologous recombination takes place using the Cre-loxP system to. . . .

In another embodiment, a vector capable of expressing CGDD or a fragment or derivative thereof may be administered to a subject to treat or prevent a. . . .

In an additional embodiment, a vector expressing the

adenovirus-based vectors are well known to those with ordinary skill in the art. Replication defective adenovirus vectors have proven to be versatile for importing genes encoding immunoregulatory proteins into intact islets in the pancreas (Csete, M.E. et al. (1995) Transplantation 27:263-268). Potentially useful adenoviral vectors are

described in U.S. Patent No. 5,707,618 to Armentano (Adenovirus

vectors for gene therapy).

hereby incorporated by reference. For adenoviral vectors, see

also Antinuzzi, P.A. et al. (1999; Annu.

which have one or more genetic abnormalities with respect to the expression of CGDD. The use of herpes simplex virus (HSV)-based vectors may be especially valuable for introducing CGDD to cells of the central nervous system, for which HSV has a tropism. The construction and packaging of herpes-based vectors are well known to those with ordinary skill in the art. A replication-competent herpes simplex virus (HSV) type I-based vector has been used to deliver a reporter gene to the eyes of primates (Liu, X. et al. (1999) Exp. Eye Res.

169:385-395). The construction of a HSV-1 virus vector has also been disclosed in detail in U.S.

For HSV vectors, see also Goins, W.F. et al. (1999; J. Virol.

73:519-532) and Xu, H. et al. (1994;

Dev. Biol. 163:152-161). The manipulation.

In another embodiment, an alphavirus (positive, single-stranded RNA virus) vector is used to deliver polynucleotides encoding CGDD to target cells. The biology of the prototypic alphavirus, Semliki Forest virus (SFV), has been studied extensively and gene transfer vectors have been based on the SFV genome (Garoff, H. and K.-J. Li (1998) Curr. Opin. Biotechnol. 9:464-469). During alphavirus RNA replication, a . . . region results in the production of a large number of CGDD-coding RNAs and the synthesis of high levels of CGDD in vector transduced cells. While alphavirus infection is typically associated with cell lysis within a few days, the ability to establish a persistent infection.

in vitro and in vivo transcription of DNA molecules encoding CGDD. Such DNA sequences may be incorporated into a wide variety of vectors with suitable RNA polymerase promoters such as T7 or SP6. Alternatively, these cDNA constructs that synthesize complementary RNA, constitutively or inducibly, can.

Many methods for introducing vectors into cells or tissues are available and equally suitable for use in vivo, in vitro, and ex vivo. For ex vivo therapy, vectors may be introduced into stem cells taken from the patient and clonally propagated for autologous transplant back into that same patient.

Means for producing specific hybridization probes for polynucleotides encoding CGDD include the cloning of polynucleotides encoding CGDD or CGDD derivatives

into vectors for the production of mRNA probes. Such vectors are known in the art, are commercially available, and may be used to synthesize RNA probes in vitro by means of.

was provided with RNA and constructed the corresponding cDNA libraries. Otherwise, cDNA was synthesized and cDNA libraries were constructed with the UNIZAP vector system (Stratagene) or SUPERScript plasmid system (Invitrogen), using the recommended procedures or similar methods known in the art (Ausubel et al.

of cDNA Clones

Plasmids obtained as described in Example I were recovered from host cells by in vivo excision using the LTNIZAP vector system (Stratagene) or by cell lysis. Plasmids were purified using

at least one of the following: a Magic or WIZARD Minipreps.

The polynucleotide sequences derived from Incyte cDNAs were validated by removing vector, linker, and poly(A) sequences and by masking ambiguous bases, using algorithms and programs based on BLAST, dynamic programming, and dinucleotide nearest.

plates, digested with CviJI cholera virus endonuclease (Molecular Biology

Research, Madison WI), and sonicated or sheared prior to religation into pUC 18 vector

(Amersham Biosciences). For shotgun

sequencing, the digested nucleotides were separated on low concentration (0.6 to 0.8%) agarose gels, fragments were excised and, . . . agar digested with Agar ACE

(Promega). Extended clones were

religated using T4 ligase (New England Biolabs, Beverly MA) into pUC 18 vector (Amersham

Biosciences), treated with Pfu DNA polymerase (Stratagene) to fill-in

restriction site overhangs, and

transfected into competent E. coli cells. Transformed cells.

by requiring a minimum Phred quality score of 15, and removed sequence alignment errors and errors resulting from improper trinucleotide of vector

sequences, chimeras, and splice

variants. An automated procedure of advanced chromosome analysis

analysed the original

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chromatogram files in the vicinity of the.

Microarray Preparation

Sequences of the present invention are used to generate array elements.

Each array element

is amplified from bacterial cells containing vectors with

cloned cDNA inserts. PCR amplification

uses primers complementary to the vector sequences flanking

the cDNA insert. Array elements are

amplified in thirty cycles of PCR from an initial quantity of 1-2 ng.

CGDD is achieved using bacterial or virus-based expression

systems. For expression of CGDD in bacteria, cDNA is subcloned into an

appropriate vector

containing an antibiotic resistance gene and an inducible promoter that

directs high levels of cDNA

transcription. Examples of such promoters include, but . . . to, the trp-lac (tac) hybrid promoter and the T5 or T7 bacteriophage promoter in conjunction with the lac operator regulatory element. Recombinant vectors are transformed into suitable bacterial hosts, e.g., BL21(DE3).

by expressing the sequences encoding CGDD at physiologically elevated levels in mammalian cell culture systems. cDNA is subcloned into a mammalian expression vector containing a strong promoter that drives high levels of cDNA expression. Vectors of choice include PCMV SPORT plasmid (Invitrogen, Carlsbad CA) and PCR3.1 plasmid (Invitrogen), both of which contain the cytomegalovirus promoter. 5-10 /tg of recombinant vector are transiently transfected into a human cell line, for example, an endothelial or hematopoietic cell line, using either liposome formulations or electroporation. . . protein provides a means to distinguish transfected cells from nontransfected cells and is a reliable predictor of cDNA expression from the recombinant vector. Marker proteins of choice include, e.g., Green Fluorescent Protein (GFP; Clontech), CD64, or a CD64-GFP fusion protein. Flow cytometry (FCM), an. . .

progression when CGDD is expressed at physiologically elevated levels in mammalian cell culture systems. cDNA is subcloned into a mammalian expression vector containing a strong promoter that drives high levels of cDNA expression. Vectors of choice include PCMV SPORT (Life Technologies, Gaithersburg, IN) and PCR 3.1 (Invitrogen, Carlsbad, CA), both of which contain the cytomegalovirus promoter. 5-10 Mg of recombinant vector are transiently transfected into a human cell line, preferably of endothelial or hematopoietic origin, using either liposome formulations or electroporation. 1-2.1-tg. . . provides a means to distinguish transfected cells from nontransfected cells and is a reliable predictor of cDNA expression from the recombinant vector.

antisense CGDD RNA [Garkawsev, I. and K. Riabowol (1997) Mol. Cell Biol. 17:2014-2019]. cDNA encoding CGDD is subcloned into the pLNCX retroviral vector to enable expression of antisense CGDD RNA. The resulting construct is transfected into the ecotropic BOSC23 virus-packaging cell line. Virus contained. . .

Alternatively, CGDD can be expressed in a mammalian cell line by transfecting the cells with a eukaryotic expression vector encoding CGDD. Eukaryotic expression vectors are commercially available, and the techniques to introduce them into cells are well known to those skilled in the art. To assay. . .

can be measured by designing an antisense sequence to the 5' end of the gene and transfecting NIH 3T3 cells with a vector transcribing this sequence. The suppression of the endogenous gene will allow transformed fibroblasts to produce clumps of cells capable of forming metastatic. . .

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L3 ANSWER 1 OF 12
ACCESSION NUMBER:
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INVENTOR(S):

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PROTEINS ASSOCIATED WITH CELL GROWTH, DIFFERENTIATION,
AND DEATH
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SK SL TJ TM TN TR TT TZ UA UG US UZ VN YU ZA ZW ZW
CH GM KE LS MW MZ SD SL SZ TZ UG ZM ZW
AM AZ BY KG KZ MD RU TJ TM
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TR
BF BJ CF CG CI CM CA GN GQ GW ML MR NE SN TD TG
WO 2002-US25465 A 20020808
US 2001-60/311,017 20010808
US 2001-60/313,070 20010817
US 2001-60/313,071 20010817
US 2001-60/314,678 20010824
US 2001-60/316,692 20010831
US 2001-60/317,913 20010907
US 2001-60/322,182 20010914
US 2001-60/340,747 20011207
US 2001-60/342,761 20011220
US 2002-60/369,129 20020329

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(FILE 'HOME' ENTERED AT 12:10:52 ON 20 MAR 2003)

FILE 'MEDLINE, CANCERLIT, BIOSIS, CONFSCI, SCISEARCH, CAPLUS, EMBASE,
USPATFULL, PCTFULL' ENTERED AT 12:11:25 ON 20 MAR 2003

L1 202 S SPAF
L2 13 S LI AND VECTOR
L3 12 DUP REM L2 (1 DUPLICATE REMOVED)

=> s l1 and DNA
L4 24 L1 AND DNA

=> dup rem l4
PROCESSING COMPLETED FOR L4
L5 19 DUP REM L4 (5 DUPLICATES REMOVED)

=> d l-19

L5 ANSWER 1 OF 19 PCTFULL COPYRIGHT 2003 Univentio
AN 2003014322 PCTFULL ED 20030303 EW 200308
TIEN PROTEINES ASSOCIATED WITH CELL GROWTH, DIFFERENTIATION, AND DEATH
TIFR PROTEINES ASSOCIEES A LA CROISSANCE, LA DIFFERENTIATION ET LA MORT
CELLULAIRES
IN AZIMZAI, Yalda, 5518 Boulder Canyon Drive, Castro Valley, CA 94552, US

[US, US];
 BARROSO, Ines, 38 Eden Street, Cambridge, Kent CB1 1EL, GB [PT, GB];
 BAUGHN, Mariah, R., 14244 Santiago Road, San Leandro, CA 94577, US [US,
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 GRAUL, Richard, C., 682-29th Avenue, San Francisco, CA 94121, US [US, US], for US only;
 GRIFFIN, Jennifer, A., 33691 Mello Way, Fremont, CA 94555, US [US, US], for US only;
 HAFALIA, April, J.A., 2227 Calle de Primavera, Santa Clara, CA 95054, US [US, US];
 ISON, Craig, H., 1242 Weathersfield Way, San Jose, CA 95118, US [US, US], for US only;
 KABLE, Amy, E., 2345 Polk Street #4, San Francisco, CA 94109, US [US, US];
 KHAN, Farrah, A., 3617 Central Road #102, Glenview, IL 60025, US [IN, US];
 LEE, Sally, 3643 26th Street, San Francisco, CA 94110, US [US, US], for US only;
 LEE, Soo Yeun, 40 Westdale Avenue, Daly City, CA 94015, US [KR, US], for US only;
 LI, Joana, X., 1264 Geneva Avenue, San Francisco, CA 94112, US [US, US], for US only;
 REDDY, Roopa, 1233 West McKinley Drive # 3, Sunnyvale, CA 94086, US [IN, US];
 RICHARDSON, Thomas, W., 616 Canyon Road #107, Redwood City, CA 94062, US [US, US], for US only;
 SPRAGUE, William, W., 611 13th Street # C, Sacramento, CA 95814, US [US, US], for US only;
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 XU, Yuming, 1739 Walnut Drive, Mountain View, CA 94040, US [US, US], for US only;
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 YUE, Henry, 826 Lois Avenue, Sunnyvale, CA 94087, US [US, US], for US only;
 YUE, Huibin, 1170 South Stelling Road, Cupertino, CA 95014, US [US, US], for US only;
 HAMLET-COX, Diana, Incyte Genomics, Inc., 3160 Porter Drive, Palo Alto, CA 94304, US [English Patent WO 2003014322 A2 20030220 W: AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ DE DK DM DZ EC EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NO NZ OM PH PL PT RO RU SD SE SG SI SK SL TJ TM TN TR TT TZ UA UG US UZ VN YU ZA ZM ZW RW (ARIPO): GH GM KE LS MW MD SD SZ TZ UG ZM ZW RW (EAP): AM AZ BY KG KZ MD RU TJ TM RW (EPO): AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE TR WO 2002-US25465 A 20020808 US 2001-60/311.017 PRAI]

NCL NCLM: 424/745.000
 NCLS: 424/757.000; 424/756.000; 514/023.000; 514/027.000; 514/053.000;
 424/755.000; 514/733.000

IC [7]
 ICM: A61K035-78
 ICS: A61K031-7C; A61K031-05
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L5 ANSWER 5 OF 19 USPATFULL
 AN 2002:301167 USPATFULL
 TI Nucleic acids, proteins, and antibodies
 IN Rosen, Craig A.; Laytonville, MD, UNITED STATES
 Ruben, Steven M.; Olney, MD, UNITED STATES
 Barash, Steven C.; Rockville, MD, UNITED STATES
 PI US 2002:168711 A1 2002:1114
 AI 2001:10117 (9)
 PRAI US 2001:764668 2000:131 (60)
 US 2000:179065P 2000:204 (60)
 US 2000:180628P 2000:0628 (60)
 US 2000:214886P 2000:0711 (60)
 US 2000:217487P 2000:0711 (60)
 US 2000:225758P 2000:0814 (60)
 US 2000:220963P 2000:0726 (60)
 US 2000:217496P 2000:0711 (60)
 US 2000:225447P 2000:0814 (60)
 US 2000:218290P 2000:0714 (60)
 US 2000:225757P 2000:0814 (60)
 US 2000:226668P 2000:0822 (60)
 US 2000:216647P 2000:0707 (60)
 US 2000:225267P 2000:0814 (60)
 US 2000:216880P 2000:0707 (60)
 US 2000:225270P 2000:0814 (60)
 US 2000:251869P 2000:1208 (60)
 US 2000:235834P 2000:0927 (60)
 US 2000:234274P 2000:0921 (60)
 US 2000:228924P 2000:0830 (60)
 US 2000:224518P 2000:0814 (60)
 US 2000:236369P 2000:0929 (60)
 US 2000:224519P 2000:0814 (60)
 US 2000:220964P 2000:0726 (60)
 US 2000:241809P 2000:1020 (60)
 US 2000:249299P 2000:1117 (60)
 US 2000:236327P 2000:0929 (60)
 US 2000:241785P 2000:1020 (60)
 US 2000:244617P 2000:1101 (60)
 US 2000:225268P 2000:0814 (60)
 US 2000:236388P 2000:0929 (60)
 US 2000:251856P 2000:1208 (60)
 US 2000:251868P 2000:1208 (60)
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 US 2000:234997P 2000:0925 (60)
 US 2000:229343P 2000:0901 (60)
 US 2000:229345P 2000:0901 (60)
 US 2000:229287P 2000:0901 (60)
 US 2000:229513P 2000:0905 (60)
 US 2000:231413P 2000:0908 (60)
 US 2000:229509P 2000:0905 (60)
 US 2000:236367P 2000:0929 (60)
 US 2000:237039P 2000:1002 (60)
 US 2000:237038P 2000:1002 (60)
 US 2000:236370P 2000:0929 (60)
 US 2000:236802P 2000:1002 (60)
 US 2000:237037P 2000:1002 (60)
 US 2000:237040P 2000:1002 (60)
 US 2000:240960P 2000:1020 (60)

US 2001-60/313.070 20010817
 US 2001-60/313.071 20010817
 US 2001-60/314.678 20010824
 US 2001-60/316.692 20010831
 US 2001-60/317.913 20010907
 US 2001-60/322.182 20010914
 US 2001-60/340.747 20011207
 US 2001-60/342.761 20011220
 US 2002-60/369.129 20020329

L5 ANSWER 2 OF 19 MEDLINE
 AN 2002741165 MEDLINE
 DN 22392627 Pubmed ID: 12504078
 TI Identification of the RT-RH/IN cleavage site of HTLV-I.
 AU Mariani Victoria L; Shuker Suzanne Beckman
 CS School of Chemistry and Biochemistry, Georgia Institute of Technology,
 Atlanta, GA 30332-0400, USA.
 SO BIOCHEMICAL AND BIOPHYSICAL RESEARCH COMMUNICATIONS, (2003 Jan 10) 300 (2)
 268-70.

CY Journal code: 0372516. ISSN: 0006-291X.
 DT Journal: Article; (JOURNAL ARTICLE)
 LA English
 FS Priority Journals
 EM 200302
 ED Entered STN: 20021231
 Last Updated on STN: 20030226
 Entered Medline: 20030225

L5 ANSWER 3 OF 19 CAPLUS COPYRIGHT 2003 ACS
 AN 2002:755245 CAPLUS
 DN 137:274175
 TI DNA, cDNA and protein sequences of spermatogenesis assocd.
 IN Kulesz-Martin, Molly F.; Liu, Yuangang
 PA USA
 SO U.S. Pat. Appl. Publ., 42 pp., Cont.-in-part of U.S. Ser. No. 777,753.
 DT Patent
 LA English
 FAN: CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2002143169	A1	20021003	US 2001-850697	20010508
US 1997-918308	B2	19970927		
US 2001-777753	A2	20010206		

L5 ANSWER 4 OF 19 USPATFULL
 AN 2002:336953 USPATFULL
 TI Medical composition for managing hormone balance
 IN Bland, Jeffrey S.; Fox Island, WA, UNITED STATES
 Liska, DeAnn J.; Gig Harbor, WA, UNITED STATES
 Tripp, Matthew, Gig Harbor, WA, UNITED STATES
 Darland, Gary K.; Gig Harbor, WA, UNITED STATES
 Lukaczer, Daniel O.; Gig Harbor, WA, UNITED STATES
 Lerman Robert, Gig Harbor, WA, UNITED STATES
 PI US 200212310 A1 20021219
 AI 2002:336953
 PRAI US 2001-265908P 20010202 (60)
 DT Utility
 FS APPLICATION
 LN: CNT 2359
 INCL INCLM: 424/745.000
 NCLM: 424/757.000; 424/756.000; 514/023.000; 514/027.000; 514/053.000;
 INCLS: 424/755.000; 514/733.000

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Center Drive, Sixteenth Floor, Newport Beach, CA 92660, US
English
Patent
WO 2002062367 AI 20020815
2002046385 PCTFULL ED 20020624 EW 200224
AE AG AU AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ
DE DK DM EZ EC EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP
KE KG KP KL LC LR LS LT LU LV MA MD MG MK MN MM NX MZ
NO NZ OM PH PL PT RO RU SD SE SG SI SK SL TJ TM TR TT TZ
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RW (EAPU): AM AZ BY CG KD MD RU TJ TM
RW (EPO): AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE TR
RW (OAPI): BF BJ CF CG CI CM GA GN GQ GW ML MR NE SN TD TG
WO 2002-US2295 A 20020125
US 2001-607265, 908 20010202
US 2002-10/056, 858 20020123
A61K035-78
ANSWER 9 OF 19 -PCTFULL COPYRIGHT 2003 Univentio
2002046385 PCTFULL ED 20020624 EW 200224
ENZYMES
ENZYMES
TANG, Y., Tom, 4230 Rawnick Court, San Jose, CA 95118, US [US, US];
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XAO, Monique G., 1189 Woodgate Drive, Camel, IN 46033, US [US, US];
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94107, US [FR, US];
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INCYTE GENOMICS, INC., 3160 Porter Drive, Palo Alto, CA 94304, US [US,
US] for all designates States except US;
TANG, Y., Tom, 4230 Rawnick Court, San Jose, CA 95118, US [US, US], for
US only;
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English
Patent
WO 2002062367 AI 20020815
2002046385 PCTFULL ED 20020624 EW 200224
AE AG AU AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ
DE DK DM EZ EC EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP
KE KG KP KL LC LR LS LT LU LV MA MD MG MK MN MM NX MZ
NO NZ OM PH PL PT RO RU SD SE SG SI SK SL TJ TM TR TT TZ
RW (ARIPO): GH GM KE LS MW MZ SD SL SZ TG UG ZM ZW
RW (EAPU): AM AZ BY CG KD MD RU TJ TM
RW (EPO): AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE TR
RW (OAPI): BF BJ CF CG CI CM GA GN GQ GW ML MR NE SN TD TG
WO 2002-US2295 A 20020125
US 2001-607265, 908 20010202
US 2002-10/056, 858 20020123
A61K035-78
ANSWER 9 OF 19 -PCTFULL COPYRIGHT 2003 Univentio
2002046385 PCTFULL ED 20020624 EW 200224
ENZYMES
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94107, US [FR, US];
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HILLMAN, Jennifer, L., 230 Monroe Drive, #17, Mountain View, CA 94040,
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INCYTE GENOMICS, INC., 3160 Porter Drive, Palo Alto, CA 94304, US [US,
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US only;

AG	STARNES, Robert, Novozymes Biotech, Inc., 1445 Drew Avenue, Davis, CA 95616, US English LA
LAF	Patent
DT	PATENT
PPI	W: WO 2002029113 A2 20020411
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AI	US 2000-09/680.598
PRAI	US 2001-60/279.526
ICM	C12Q001-68 20010327
	ANSWER 11 OF 19 PCTFULL COPYRIGHT 2003 Univentio
L5	2002018434 PCTFULL ED 20020705 EW 200210
AN	PROTEINS RELATED TO SCHIZOPHRENIA AND USES THEREOF
TITION	PROTEINES LIEES A LA SCHIZOPHRENIE ET UTILISATIONS DE CELLULES-CI
TITIFR	ST.GEORGE-HYSLOP, Peter, H., 210 Richview Avenue, Toronto, Ontario M5P 3G3, CA;
IN	FRASER, Paul, E., 611 Windermere Avenue, Toronto, Ontario M6S 3L9, CA THE GOVERNING COUNCIL OF THE UNIVERSITY OF TORONTO, Simcoe Hall Room 131S, 27 King's College Circle, Toronto, Ontario M5S 1A1, CA (CA, CA) RAE, Patricia, A., Sim & McBurney, 330 University Avenue, 6th Floor, Toronto, Ontario M5G 1R7, CA
AG	English PATENT
LAF	English
LA	Patent
DT	PATENT
PPI	W: WO 2002018434 A2 20020307
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	RW (ARIPO): GH GM KE LS MM MZ SD SL SZ Tz UG ZW RW (EPO): AM AZ BY CY KZ MD RU TJ TM RW (EP0): AT BE CH CG DE DK ES FI FR GB GR IE IT LU MC NL PT SE TR RW (OAPI): BF BJ CF CG CI CM GA GN GQ GW ML MR NE SN TD TG WO 2001-CA1243 A 20010831
AI	US 2000-60/229.889
PRAI	US 2000-60/229.889
ICM	C07K014-47 20000901
	ANSWER 12 OF 19 PCTFULL COPYRIGHT 2003 Univentio
L5	2002006339 PCTFULL ED 20020814
AN	NOVEL PROTEINS AND NUCLEIC ACIDS ENCODING SAME
TITION	NOUVELLES PROTEINES ET ACIDES NUCLEIQUES LES CODANT
TITIFR	SPADERNA, Steven, K.;
IN	TCHERNEV, Velizar; Liu, Xiaohong; SHENOY, Suresh; SPYTEK, Kimberly; ZERHUSEN, Bryan; PATIRAJAN, Meera; TAUPIER, Raymond, J.; RASTELLI, Luca; GROSSE, William, M.; SZEKERES, Edward, S.; ALSOBROOK, John, II; LEPLY, Denise, M.

PA	SHEN, Lei; BURGESS, Catherine, E.; SHIMKETS, Richard, A.; PADIGARU, Muralidhara CURAGEN CORPORATION; SPADERNA, Steven, K.; TCHERNEV, Velizar; LIU, Xiaohong; SHENOY, Suresh; SPYTEK, Kimberly; ZERHUSEN, Bryan; PATTURAJAN, Meera; TAUPIER, Raymond, J.; RASTELLI, Luca; GROSSE, William, M.; SZEKERES, Edward, S.; ALSOBROOK, John, II; LEPLEY, Denise, M.; SHEN, Lei; BURGESS, Catherine, E.; SHIMKETS, Richard, A.; PADIGARU, Muralidhara Patent WO 2002006339	AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ DE DK DM DZ EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MA MD MG MN MW MX MY NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR TT TZ UA UG US UZ VN YU ZA ZW CH CM KE LS MW MZ SD SL SZ TZ UG ZW AM AZ BY BG KZ MD RU TJ TM AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE TR BF BU CF CG CI CM GN GW ML MR NE SN TD TG	A2 20020124	US 2000-60/179,065 US 2000-60/180,628 US 2000-60/184,664 US 2000-60/186,350 US 2000-60/189,874 US 2000-60/190,076 US 2000-60/198,123 US 2000-60/205,515 US 2000-60/209,467 US 2000-60/214,886 US 2000-60/215,135 US 2000-60/216,647 US 2000-60/216,880 US 2000-60/217,487 US 2000-60/217,496 US 2000-60/218,290 US 2000-60/220,963 US 2000-60/220,964 US 2000-60/225,757 US 2000-60/225,270 US 2000-60/225,447 US 2000-60/225,457 US 2000-60/225,758 US 2000-60/225,758 US 2000-60/225,268 US 2000-60/224,518 US 2000-60/224,519 US 2000-60/225,759 US 2000-60/225,213 US 2000-60/225,266 US 2000-60/225,214 US 2000-60/226,279, US 2000-60/226,868 US 2000-60/227,182 US 2000-60/226,681 US 2000-60/227,009 US 2000-60/228,924 US 2000-60/229,344 US 2000-60/229,343 US 2000-60/229,287 US 2000-60/229,345 US 2000-60/229,513 US 2000-60/229,509 US 2000-60/230,438 US 2000-60/230,437 US 2000-60/231,413 US 2000-60/232,080 US 2000-60/231,414 US 2000-60/231,244 US 2000-60/232,081 US 2000-60/231,242 US 2000-60/231,243 US 2000-60/231,968 US 2000-60/232,401 US 2000-60/232,399 US 2000-60/232,400 US 2000-60/232,397 US 2000-60/233,063 US 2000-60/233,064 US 2000-60/233,065 US 2000-60/232,398 US 2000-60/234,223 US 2000-60/234,274 US 2000-60/234,997	WO 2001-US1312 US 2000-60/179,065 US 2000-60/180,628 US 2000-60/184,664 US 2000-60/186,350 US 2000-60/189,874 US 2000-60/190,076 US 2000-60/198,123 US 2000-60/205,515 US 2000-60/209,467 US 2000-60/214,886 US 2000-60/215,135 US 2000-60/216,647 US 2000-60/216,880 US 2000-60/217,487 US 2000-60/217,496 US 2000-60/218,290 US 2000-60/220,963 US 2000-60/220,964 US 2000-60/225,757 US 2000-60/225,270 US 2000-60/225,447 US 2000-60/225,457 US 2000-60/225,758 US 2000-60/225,758 US 2000-60/225,268 US 2000-60/224,518 US 2000-60/224,519 US 2000-60/225,759 US 2000-60/225,213 US 2000-60/225,266 US 2000-60/225,214 US 2000-60/226,279, US 2000-60/226,868 US 2000-60/227,182 US 2000-60/226,681 US 2000-60/227,009 US 2000-60/228,924 US 2000-60/229,344 US 2000-60/229,343 US 2000-60/229,287 US 2000-60/229,345 US 2000-60/229,513 US 2000-60/229,509 US 2000-60/230,438 US 2000-60/230,437 US 2000-60/231,413 US 2000-60/232,080 US 2000-60/231,414 US 2000-60/231,244 US 2000-60/232,081 US 2000-60/231,242 US 2000-60/231,243 US 2000-60/231,968 US 2000-60/232,401 US 2000-60/232,399 US 2000-60/232,400 US 2000-60/232,397 US 2000-60/233,063 US 2000-60/233,064 US 2000-60/233,065 US 2000-60/232,398 US 2000-60/234,223 US 2000-60/234,274 US 2000-60/234,997	PT SE TR BF BJ CF CG CI CM GN GW ML MR NE SN TD TG	WO 2001-US1312 US 2000-60/179,065 US 2000-60/180,628 US 2000-60/184,664 US 2000-60/186,350 US 2000-60/189,874 US 2000-60/190,076 US 2000-60/198,123 US 2000-60/205,515 US 2000-60/209,467 US 2000-60/214,886 US 2000-60/215,135 US 2000-60/216,647 US 2000-60/216,880 US 2000-60/217,487 US 2000-60/217,496 US 2000-60/218,290 US 2000-60/220,963 US 2000-60/220,964 US 2000-60/225,757 US 2000-60/225,270 US 2000-60/225,447 US 2000-60/225,457 US 2000-60/225,758 US 2000-60/225,758 US 2000-60/225,268 US 2000-60/224,518 US 2000-60/224,519 US 2000-60/225,759 US 2000-60/225,213 US 2000-60/225,266 US 2000-60/225,214 US 2000-60/226,279, US 2000-60/226,868 US 2000-60/227,182 US 2000-60/226,681 US 2000-60/227,009 US 2000-60/228,924 US 2000-60/229,344 US 2000-60/229,343 US 2000-60/229,287 US 2000-60/229,345 US 2000-60/229,513 US 2000-60/229,509 US 2000-60/230,438 US 2000-60/230,437 US 2000-60/231,413 US 2000-60/232,080 US 2000-60/231,414 US 2000-60/231,244 US 2000-60/232,081 US 2000-60/231,242 US 2000-60/231,243 US 2000-60/231,968 US 2000-60/232,401 US 2000-60/232,399 US 2000-60/232,400 US 2000-60/232,397 US 2000-60/233,063 US 2000-60/233,064 US 2000-60/233,065 US 2000-60/232,398 US 2000-60/234,223 US 200
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PI WO 2000072854 A1 20001207
DS AE AG AL AM AT AU AZ BA BB BG BR BY CA CH CN CR CU CZ DE DK
DM DZ EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP
KR KZ LC LR LS LT LU LV MA MD MG MK MN MX MY NZ NO NZ PL
PT RO RU SD SE SG SI SK SL TJ TM TR TT TZ UA UG US UZ VN YU
ZA ZW GM KE LS MW MZ SD SL SZ T2 UG ZW AM AZ BY KG KZ MD
RU TJ TM AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT
SE BF BJ CF CG CI CM GN GW ML MR NE SN TD TG

AI WO 2000-US15196 A 20000602
PRAI US 1999-60/137.080 19990602

L5 ANSWER 17 OF 19 PCTFULL COPYRIGHT 2003 Univentio
AN 200006069 PCTFULL ED 20020515
TIEN A PRESENILIN ASSOCIATED MEMBRANE PROTEIN AND USES THEREOF
TIFR PROTEINE MEMBRANAIRE ASSOCIEE A LA PRESENILINE ET SES UTILISATIONS
IN ST. GEORGE-HYSLUP, Peter, H.;
FRASER, Paul, E.
PA THE GOVERNING COUNCIL OF THE UNIVERSITY OF TORONTO
LA English
DT Patent

DS WO 2000060069 A1 20001012
W: AE AG AL AM AT AU AZ BA BB BG BR BY CA CH CN CR CU CZ DE DK
DM DZ EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP
KR KZ LC LR LS LT LU LV MA MD MG MK MN MX MY NZ NO NZ PL
PT RO RU SD SE SG SI SK SL TJ TM TR TT TZ UA UG US UZ VN YU
ZA ZW GM KE LS MW SD SL SZ T2 UG ZW AM AZ BY KG KZ MD RU TJ TM
AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE BF BJ
CF CG CI CM CA CN GM ML MR NE SN TD TG

AI WO 2000-CA354
PRAI US 1999-60/127.452 19990401
ICM US 1999-60/173.826 19991230
ICS C07K014-705; A01K067-027; C12N005-10; C120001-68; G01N033-50

L5 ANSWER 18 OF 19 MEDLINE
AN 2000200628 MEDLINE
DN 20200628 Pubmed ID: 10734318
TI SPAP, a new AAA-protein specific to early spermatogenesis and
malignant conversion.
AU Liu Y; Black J; Kiesel N; Kulesz-Martin M F
CS Program of Biochemistry and Department of Pharmacology and Therapeutics,
Roswell Park Cancer Institute, Elm & Carlton Streets, Buffalo, New York,
NY 14263, USA.
NC C01C031101 (NCI)
C01C031101 (NCI)
SO ONCOGENE, (2000 Mar 16) 19 (12) 1579-88.
CY Journal code: 8711562. ISSN: 0950-9232.
DT JOURNAL: United Kingdom
LA English
FS Priority Journals
OS GENBANK-AF049099
EM 200004
ED Entered STN: 20000505
Last Updated on STN: 20000505
Entered Medline: 20000421

L5 ANSWER 19 OF 19 MEDLINE
AN 94368094 MEDLINE
DN 94368094 Pubmed ID: 8085823
TI Genes involved in self-protection against the lantibiotic subtilin
produced by Bacillus subtilis ATCC 6633.
AU Klein C; Enian K D
CS Institute for Microbiology, Biozentrum der Johann Wolfgang
Goethe-Universität, Frankfurt am Main, Federal Republic of Germany.

SO APPLIED AND ENVIRONMENTAL MICROBIOLOGY, (1994 Aug) 60 (8) 2793-801.
CY Journal code: 7605801. ISSN: 0099-2240.
DT United States
LA English
FS Priority Journals
OS GENBANK-U09819
EM 199410
ED Entered STN: 19941021
Last Updated on STN: 19941021
Entered Medline: 19941011

=> d his
(FILE 'HOME' ENTERED AT 12:10:52 ON 20 MAR 2003)
FILE 'MEDLINE, CANCERLIT, BIOSIS, CONFSCI, SCISEARCH, CAPLUS, EMBASE,
USPATFULL, PCTFULL' ENTERED AT 12:11:25 ON 20 MAR 2003
202 S SPAP
L1 13 S L1 AND VECTOR
L2 12 DUP REM L2 (1 DUPLICATE REMOVED)
L3 24 S L1 AND DNA
L4 19 DUP REM L4 (5 DUPLICATES REMOVED)
L5
=> s l1 and CDNA
L6 13 L1 AND CDNA
=> dup rem l6
PROCESSING COMPLETED FOR L6
L7 12 DUP REM L6 (1 DUPLICATE REMOVED)
=> d 1-12

L7 ANSWER 1 OF 12 PCTFULL COPYRIGHT 2003 Univentio
AN 2003014322 PCTFULL ED 20030303 EW 200308
TIEN PROTEINS ASSOCIATED WITH CELL GROWTH, DIFFERENTIATION, AND DEATH
TIFR PROTEINES ASSOCIEES A LA CROISSANCE, LA DIFFERENTIATION ET LA MORT
CELLULAIRES
IN AZIMZAI, Valda, 5518 Boulder Canyon Drive, Castro Valley, CA 94552, US
[US, US];
BARROSO, Ines, 38 Eden Street, Cambridge, Kent CB1 1EL, GB [PT, GB];
BAUGHN, Mariah, R., 14244 Santiago Road, San Leandro, CA 94577, US [US,
US];
BECHA, Shanya, D., 21062 Gary Drive # 117, Castro Valley, CA 94546, US
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US [AU, US];
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US];
GORVAD, Ann, E., 369 Marie Common, Livermore, CA 94550, US [US, US];
GRAUB, Richard, C., 682-29th Avenue, San Francisco, CA 94121, US [US,
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KHAN, Farrah, A., 3617 Central Road #102, Glenview, IL 60025, US [IN, US];
LEE, Sally, 3643 26th Street, San Francisco, CA 94110, US [US, US];
LEE, Soo Yeun, 40 Westdale Avenue, Daly City, CA 94015, US [KR, US];
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REDDY, Roopa, 1233 West McKinley Drive # 3, Sunnyvale, CA 94086, US [IN, US];
RICHARDSON, Thomas, W., 616 Canyon Road #107, Redwood City, CA 94062, US [US, US];
SPRAGUE, William, W., 611 13th Street # C, Sacramento, CA 95814, US [US, US];
TANG, Y. Tom, 4230 Ranwick Court, San Jose, CA 95118, US [US, US];
WARREN, Bridget, A., 1810 S. El Camino Real #B103, Encinitas, CA 94024, US [US, US];
XU, Yuming, 1739 Walnut Drive, Mountain View, CA 94040, US [US, US];
YAO, Monique, G., 1189 Woodgate Drive, Carmel, IN 46033, US [US, US];
YUE, Henry, 826 Lois Avenue, Sunnyvale, CA 94087, US [US, US];
YUE, Huibin, 1170 South Stelling Road, Cupertino, CA 95014, US [US, US];
INCYTE GENOMICS, INC., 3160 Porter Drive, Palo Alto, CA 94304, US [US, US];
for all designates States except US;
AZIMZAI, Valda, 5518 Boulder Canyon Drive, Castro Valley, CA 94552, US [US, US];
for US only;
BARROSO, Ines, 38 Eden Street, Cambridge, Kent CB1 1EL, GB [PT, GB], for US only;
BAUGHN, Mariah, R., 14244 Santiago Road, San Leandro, CA 94577, US [US, US];
for US only;
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for US only;
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HAFALIA, April, J.A., 2227 Calle de Primavera, Santa Clara, CA 95054, US [US, US];
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ISON, Craig, H., 1242 Weathersfield Way, San Jose, CA 95118, US [US, US];
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KABLE, Amy, E., 2345 Polk Street #4, San Francisco, CA 94109, US [US, US];
for US only;
KHAN, Farrah, A., 3617 Central Road #102, Glenview, IL 60025, US [IN, US];
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LEE, Sally, 3643 26th Street, San Francisco, CA 94110, US [US, US];
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for US only;

US only;
LI, Joana, X., 1264 Geneva Avenue, San Francisco, CA 94112, US [US, US];
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REDDY, Roopa, 1233 West McKinley Drive # 3, Sunnyvale, CA 94086, US [IN, US];
for US only;
RICHARDSON, Thomas, W., 616 Canyon Road #107, Redwood City, CA 94062, US [US, US];
for US only;
SPRAGUE, William, W., 611 13th Street # C, Sacramento, CA 95814, US [US, US];
for US only;
SWARNAKAR, Anita, 8 Locksley Avenue #50, San Francisco, CA 94122, US [CA, US];
for US only;
TANG, Y. Tom, 4230 Ranwick Court, San Jose, CA 95118, US [US, US];
for US only;
WARREN, Bridget, A., 1810 S. El Camino Real #B103, Encinitas, CA 94024, US [US, US];
for US only;
XU, Yuming, 1739 Walnut Drive, Mountain View, CA 94040, US [US, US];
for US only;
YAO, Monique, G., 1189 Woodgate Drive, Carmel, IN 46033, US [US, US];
for US only;
YUE, Henry, 826 Lois Avenue, Sunnyvale, CA 94087, US [US, US];
for US only;
YUE, Huibin, 1170 South Stelling Road, Cupertino, CA 95014, US [US, US];
for US only;
HAMLET-COX, Diana, Incyte Genomics, Inc., 3160 Porter Drive, Palo Alto, CA 94304, US [US, US];
English
LAF Patent
LA English
DT Patent
PI WO 20030014322 A2 20030220
W: AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ DE DK DM DZ EC EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NO NZ OM PH PL PT RO RU SD SE SG SI SK SL TJ TM TN TR TT TZ UA US UZ VN YU ZA ZW ZW
RW (ARIPO): GH GM KE LS MW MZ SD SL SZ TZ UG ZM ZW
RW (EAP): AM AZ BY KG KZ MD RU TJ TM
RW (EPO): AT BE CH CY DE DK ES FI FR GB GR IE IT LJ MC NL PT SE TR
RW (OAPI): BF BJ CF CG CI CM GN GQ GW ML MR NE SN TD TG
A 20020808
AI WO 2002-US25465 A 20020808
PRAI US 2001-60/311.017 20010808
US 2001-60/313.070 20010817
US 2001-60/313.071 20010817
US 2001-60/314.678 20010824
US 2001-60/316.692 20010831
US 2001-60/317.913 20010907
US 2001-60/322.182 20010914
US 2001-60/340.747 20011207
US 2001-60/342.761 20011220
US 2002-60/369.129 20020329
L7 ANSWER 2 OF 12 CAPLUS COPYRIGHT 2003 ACS DUPLICATE 1
AN 2002:755245 CAPLUS
DN 137:274175
TI DNA, cDNA and protein sequences of spermatogenesis assocd.
IN Kulesz-Martin, Molly F.; Liu, Yuangang
PA USA Pat. Appl. Publ., 42 pp., Cont.-in-part of U.S. Ser. No. 777,753.
SO CODEN: USXXCO
DT Patent
LA English
FAN CNT 1
PI US 2002143169 A1 20021003 APPLICATION NO. DATE
US 2001-850697 20010508

PRAI US 1997-938308 B2 19970927
US 2001-777753 A2 20010206

L7 ANSWER 3 OF 12 USPATFULL
AN 2002:301167 USPATFULL
TI Nucleic acids, proteins, and antibodies
IN Ruben, Craig A., Laytonville, MD, UNITED STATES
Barash, Steven C., Rockville, MD, UNITED STATES
PRAI US 2001-764868 A1 20010117 (9)
US 2000-179065P 20000131 (60)
US 2000-180628P 20000204 (60)
US 2000-214886P 20000628 (60)
US 2000-217487P 20000711 (60)
US 2000-225758P 20000814 (60)
US 2000-220963P 20000726 (60)
US 2000-217496P 20000711 (60)
US 2000-225447P 20000814 (60)
US 2000-218290P 20000714 (60)
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US 2000-236369P 20000929 (60)
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US 2000-220964P 20000726 (60)
US 2000-241809P 20001020 (60)
US 2000-249299P 20001117 (60)
US 2000-236327P 20000929 (60)
US 2000-241785P 20001020 (60)
US 2000-244617P 20001101 (60)
US 2000-225268P 20000814 (60)
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US 2000-237039P 20001002 (60)
US 2000-237038P 20001002 (60)
US 2000-236370P 20000929 (60)
US 2000-236802P 20001002 (60)
US 2000-237034P 20001002 (60)
US 2000-237040P 20001002 (60)
US 2000-240960P 20001020 (60)
US 2000-239935P 20001013 (60)
DT Utility
FS APPLICATION
LN.CNT 31967
INCL INCLM: 435/069.100

INCLM: 435/325.000; 435/320.100; 435/183.000; 530/350.000; 536/023.100
NCLM: 435/069.100
NCLS: 435/325.000; 435/320.100; 435/183.000; 530/350.000; 536/023.100

IC [7]
ICM: C12P021-02
ICS: C12N005-06; C07H021-04; C12N009-00; C07K014-435
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 4 OF 12 USPATFULL
AN 2002:112541 USPATFULL
TI Proteins related to schizophrenia and uses thereof
IN St. George-Hyslop, Peter H., Toronto, CANADA
Fraser, Paul E., Toronto, CANADA
PA The Governing Council of the University of Toronto (non-U.S. corporation)
PI US 2002058276 A1 20020516
AI US 2001-945258 A1 20010831 (9)
PRAI US 2000-229889P 20000901 (60)
DT Utility
FS APPLICATION
LN.CNT 2309
INCL INCLM: 435/006.000
INCLM: 424/009.200; 803/003.000
NCLM: 435/006.000
NCLS: 424/009.200; 803/003.000

IC [7]
ICM: C12Q001-68
ICS: A61K049-00; A01K057-00
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 5 OF 12 PCTFULL COPYRIGHT 2003 Univentio
AN 2002099062 PCTFULL ED 20021218 EW 200250
TIEN NOVEL ANTIBODIES THAT BIND TO ANTIGENIC POLYPEPTIDES, NUCLEIC ACIDS
ENCODING THE ANTIGENS, AND METHODS OF USE
TIFR NOUVEAUX ANTICORPS SE FIXANT A DES POLYPEPTIDES ANTIGENIQUES, ACIDES
NUCLEIQUES CODANT LES ANTIGENES ET MODES D'UTILISATION
IN ANDERSON, David, W., 85 Montoya Drive, Branford, CT 06405, US [US, US];
ZERHUSEN, Bryan, D., 337 Monticello Drive, Branford, CT 06405, US [US, US];
LI, Li, 56 Jerimoth Drive, Branford, CT 06405, US [CN, US];
ZHONG, Mei, 45 Harrison Avenue, Apartment 1B, Branford, CT 06405, US [CA, US];
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GERLACH, Valerie, L., 18 Rock Pasture Road, Branford, CT 06405, US [US, US];
SHIMKETS, Richard, A., 5 Indian Meadows Drive, Guilford, CT 06437, US [US, US];
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PATTURAJAN, Meera, 45 Harrison Avenue, Apartment 1C, Branford, CT 06405, US [IN, US];
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MACDOUGALL, John, R., 117 Russell Street, Hamden, CT 06517, US [CA, US];
TAUPIER, Raymond, J., Jr., 34 Pardee Place Extension, East Haven, CT 06512, US [US, US];
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 BOLDGOG, Ferenc, L., 1687 Hartford Turnpike, North Haven, CT 06473, US
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AG	ELRIFI, Ivor. R., Mintz, Levin, Cohn, Ferris, Glosky, and Popeo, P.,
C., One Financial Center, Boston, MA 02111. US	
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ENZYMES
TIEN
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IN
TANG, Y. Y., Tom, 4230 Ranwick Court, San Jose, CA 95118, US [US, US];
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 INCYTE GENOMICS, INC., 3160 Porter Drive, Palo Alto, CA 94304, US [US,
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 TANG, Y., Tom, 4230 Ranwick Court, San Jose, CA 95118, US [US, US], for
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 YUE, Henry, 826 Lois Avenue, Sunnyvale, CA 94087, US [US, US], for US
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PA

LAF English
 LA English
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 US 2000-60/254,312 20001208
 US 2000-60/255,773 20001214
 US 2000-60/256,188 20001215
 US 2000-60/255,940 20001215
 US 2000-60/257,488 20001221
 US 2001-60/262,839 20010119
 US 2001-60/264,402 20010126
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 L7 ANSWER 7 OF 12 PCTFULL COPYRIGHT 2003 Univentio
 AN 2002029113 PCTFULL ED 20020627 EW 200215
 TIEN METHODS FOR MONITORING MULTIPLE GENE EXPRESSION
 TIFR METHODES DE SURVEILLANCE DE L'EXPRESSION GENETIQUE MULTIPLE
 IN BERKA, Randy, 3609 Modoc, Davis, CA 95616, US;
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 AG STARNES, Robert, Novozymes Biotech, Inc., 1445 Drew Avenue, Davis, CA
 95616, US
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 LA English
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 US 2001-60/279,526 20010327
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 L7 ANSWER 8 OF 12 PCTFULL COPYRIGHT 2003 Univentio
 AN 2002018434 PCTFULL ED 20020705 EW 200210
 TIEN PROTEINES RELATED TO SCHIZOPHRENIA AND USES THEREOF
 TIFR PROTEINES LIEES A LA SCHIZOPHRENIE ET UTILISATIONS DE CELLES-CI
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 LAF English
 LA English

AG

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AI WO 2001-CA1243 A 20010831
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L7 ANSWER 9 OF 12 PCTFULL COPYRIGHT 2003 Univentio
AN 200206339 PCTFULL ED 20020814
TIEN NOVEL PROTEINS AND NUCLEIC ACIDS ENCODING SAME
TIFF NOUVELLES PROTEINES ET ACIDES NUCLEIQUES LES CODANT
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SHEN, Lei;
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L7 ANSWER 10 OF 12 PCTFULL COPYRIGHT 2003 Univentio
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TIEN NUCLEIC ACIDS, PROTEINS AND ANTIBODIES
TIFF ACIDES NUCLEIQUES, PROTEINES ET ANTICORPS
IN ROSEN, Craig, A.;
BARASH, Steven, C.;
RUBEN, Steven, M.;
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ROSEN, Craig, A.;
BARASH, Steven, C.;
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AG1K048-00
C12N005-00

ANSWER 11 OF 12 PCTFULL COPYRIGHT 2003 Univentio
2000060069 PCTFULL ED 20020515
A PRESENILIN ASSOCIATED MEMBRANE PROTEIN AND USES THEREOF
PROTEINE MEMBRANAIRE ASSOCIEE A LA PRESENILINE ET SES UTILISATIONS
ST. GEORGE-HYSLOP, Peter, H.;
FRASER, Paul, E.
THE GOVERNING COUNCIL OF THE UNIVERSITY OF TORONTO
English
Patent
WO 2000060069 A1 20001012
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US 1999-60/127, 452
US 1999-60/173, 826
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C07K014-705; A01K067-027; C12N005-10; C12Q001-68; G01N033-50

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L7 ANSWER 12 OF 12 CAPLUS COPYRIGHT 2003 ACS
AN 2000:242308 CAPLUS
DN 133:29064
TI SPAP, a new AAA-protein specific to early spermatogenesis and
   malignant conversion
AU Liu, Yungang, Black, Jennifer; Kisiel, Nicholas; Kulesz-Martin, Molly F.
CS Program of Biochemistry and Department of Pharmacology and Therapeutics,
   Roswell Park Cancer Institute, Buffalo, NY, 14263, USA
SO Oncogene (2000), 19(12), 1579-1588
   CODEN: ONCNE5; ISSN: 0950-9232
PB Nature Publishing Group
DT Journal
LA English
RE.CNT 44 THERE ARE 44 CITED REFERENCES AVAILABLE FOR THIS RECORD
          ALL CITATIONS AVAILABLE IN THE RE FORMAT

```

```

=> d his

```

```

(FILE 'HOME' ENTERED AT 12:10:52 ON 20 MAR 2003)
FILE 'MEDLINE, CANCERLIT, BIOSIS, CONFSCI, SCISEARCH, CAPLUS, EMBASE,
USPATFULL, PCTFULL' ENTERED AT 12:11:25 ON 20 MAR 2003
L1 202 S SPAP
L2 13 S L1 AND VECTOR
L3 12 DUP REM L2 (1 DUPLICATE REMOVED)
L4 24 S L1 AND DNA
L5 19 DUP REM L4 (5 DUPLICATES REMOVED)
L6 13 S L1 AND CDNA
L7 12 DUP REM L6 (1 DUPLICATE REMOVED)
=> s atcc (a) 98558
L8 0 ATCC (A) 98558
=>

```

```

---Logging off of STN---

```

```

=>
Executing the logoff script...

```

```

=> LOG Y
COST IN U.S. DOLLARS          SINCE FILE          TOTAL
ENTRY                          60.59          SESSION
FULL ESTIMATED COST          60.80
STN INTERNATIONAL LOGOFF AT 12:23:53 ON 20 MAR 2003

```